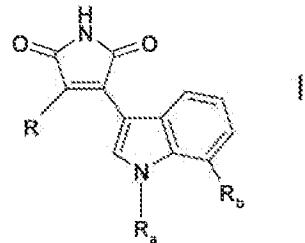


Amendments to the claims.

This listing of claims will replace all prior versions and listings of claims in the application.

Listing of Claims.

1. (currently amended) A compound of formula I

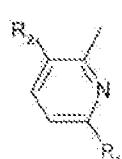


wherein

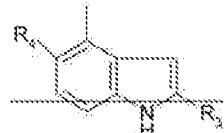
$R_a$  is H;  $C_{1-4}$ alkyl; or  $C_{1-4}$ alkyl substituted by OH,  $NH_2$ ,  $NHC_{1-4}$ alkyl or  $N(di-C_{1-4}$ alkyl) $_2$ ;

$R_b$  is H; halogen;  $C_{1-6}$ alkyl; or  $C_{1-6}$ alkoxy, and

$R$  is a radical of formula (a) or (b)



(a)



(b)

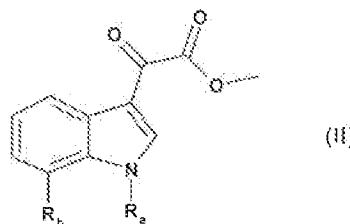
wherein

each of  $R_1$  is piperazine and  $R_3$  is a heterocyclic residue; or a radical of formula  $\alpha$

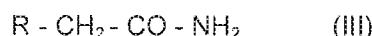


wherein X is a direct bond, O, S or NR<sub>14</sub>, wherein R<sub>14</sub> is H or C<sub>1-4</sub>alkyl;  
R<sub>6</sub> is C<sub>1-4</sub>alkylene or C<sub>1-4</sub>alkylene wherein one CH<sub>2</sub> is replaced by CR<sub>x</sub>R<sub>y</sub>, wherein one of R<sub>x</sub> and R<sub>y</sub> is H and the other is CH<sub>3</sub>, each of R<sub>x</sub> and R<sub>y</sub> is CH<sub>3</sub> or R<sub>x</sub> and R<sub>y</sub> form together CH<sub>2</sub>CH<sub>2</sub>;  
Y is bound to the terminal carbon atom and is selected from OH, NR<sub>12</sub>R<sub>13</sub> wherein each of R<sub>12</sub> and R<sub>13</sub>, independently, is H, C<sub>3-6</sub>cycloalkyl, C<sub>3-6</sub>cycloalkyl-C<sub>1-4</sub>alkyl, aryl, aryl-C<sub>1-4</sub>alkyl, heteroaryl-C<sub>1-4</sub>alkyl, C<sub>2-6</sub>alkenyl or C<sub>1-4</sub>alkyl optionally substituted on the terminal carbon atom by OH, halogen, C<sub>1-4</sub>alkoxy or NR<sub>14</sub>R<sub>15</sub> wherein each of R<sub>14</sub> and R<sub>15</sub>, independently, is H, C<sub>1-4</sub>alkyl, C<sub>3-6</sub>cycloalkyl, C<sub>3-6</sub>cycloalkyl-C<sub>1-4</sub>alkyl, aryl-C<sub>1-4</sub>alkyl, or R<sub>12</sub> and R<sub>13</sub> form together with the nitrogen atom to which they are bound a heterocyclic residue; and  
each of R<sub>2</sub> and R<sub>4</sub>, independently, is H; halogen; C<sub>1-4</sub>alkyl; C<sub>1-4</sub>alkoxy; CF<sub>3</sub>; nitrile; nitro or amino,  
or a salt thereof.

2. (original) A compound according to claim 1 wherein  $R_a$  is H, methyl, ethyl, or isopropyl, or a salt thereof.
3. (currently amended) A compound according to claim 1 or 2 wherein  $R_b$  is H, Cl, methyl or ethyl, or a salt thereof.
4. (currently amended) A compound according to any one of claims 1 to 3 wherein  $R_1$  is a heterocyclic residue, e.g. a piperazinyl, optionally substituted on a ring nitrogen or on a ring carbon, e.g. 4-methyl-piperazin-1-yl, or 4,7-diaza-spiro[2.5]oct-7-yl; or a radical of formula (a) wherein X is a direct bond,  $R_6$  is  $CH_2$  and Y is  $NR_{12}R_{13}$  wherein each of  $R_{12}$  and  $R_{13}$ , independently, is H,  $C_{3-6}$ cycloalkyl- $C_{1-4}$ alkyl,  $C_{2-6}$ alkenyl or  $C_{1-4}$ alkyl optionally substituted on the terminal carbon atom by OH, halogen,  $C_{1-4}$ alkoxy or  $NR_{14}R_{15}$  wherein each of  $R_{14}$  and  $R_{15}$ , independently, is H or  $C_{1-4}$ alkyl; or  $R_{12}$  and  $R_{13}$  form together with the nitrogen atom to which they are bound a heterocyclic residue e.g. a piperazinyl, or a salt thereof.
5. (currently amended) A compound according to any one of claims 1 to 4 wherein  $R_2$  and/or  $R_4$  is H, Cl, F,  $CF_3$ , nitrile, nitro or amino, or a salt thereof.
6. (previously presented): A process for the preparation of a compound of formula I according to claim 1, which process comprises reacting a compound of formula II



wherein  $R_a$  and  $R_b$  are as defined in claim 1,  
with a compound of formula III



wherein R is as defined in claim 1,  
and, where required, converting the resulting compound of formula I obtained in free form to a salt form or vice versa, as appropriate.

7. (currently amended) A compound of formula I according to any one of claims 1 to 5, in free form or in a pharmaceutically acceptable salt form for use as a pharmaceutical.
8. (currently amended) A pharmaceutical composition comprising a compound of formula I according to any one of claims 1 to 5, in free form or in a pharmaceutically acceptable salt form, in association with a pharmaceutically acceptable diluent or carrier therefor.
9. (cancelled).

10 (currently amended): A method for preventing or treating acute or chronic transplant rejection disorders or diseases mediated by T-lymphocytes and/or PKC or GSK-3 $\beta$  in a subject in need of such treatment, which method comprises administering to said subject an effective amount of a compound of formula I according to any one of claims 1 to 5 or a pharmaceutically acceptable salt thereof.

11 (new) A method for preventing or treating T-cell mediated inflammatory or autoimmune diseases, in a subject in need of such treatment, which method comprises administering to said subject an effective amount of a compound of formula I of claim 1 or a pharmaceutically acceptable salt thereof.